Vitamin AD3E pro injectione, solution for injection for horses, cattle, pigs, and dogs

Not authorised

- Retinol palmitate
- COLECALCIFEROL CONCENTRATE (OILY FORM)
- DL-ALPHA TOCOPHEROL ACETATE

Product identification

Medicine name:
Vitamin AD3E pro injectione, solution for injection for horses, cattle, pigs, and dogs
Belavit AD3E, raztopina za injiciranje za konje, govedo, prašiče in pse

Active substance:
Retinol palmitate
COLECALCIFEROL CONCENTRATE (OILY FORM)
DL-ALPHA TOCOPHEROL ACETATE

Target species:
Cattle
Dog
Horse
Pig

Route of administration:
Subcutaneous use
Intramuscular use

Product details

Active substance and strength:

Retinol palmitate
176.46 milligram(s) / 1.00 millilitre(s)

COLECALCIFEROL CONCENTRATE (OILY FORM)
100.00 milligram(s) / 1.00 millilitre(s)

DL-ALPHA TOCOPHEROL ACETATE
50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - Milk. 120 hour
  - Meat and offal. 259 day
- **Horse**
  - Meat and offal. 250 day
  - Milk. no withdrawal period

Not authorised for use in horses producing milk for human consumption.

- **Pig**
  - Meat and offal. 194 day

Intramuscular use:

- **Cattle**
  - Milk. 120 hour

- Meat and offal. 259 day

- **Horse**
  - Meat and offal. 250 day
  - Milk. no withdrawal period

Not authorised for use in horses producing milk for human consumption.

- **Pig**
  - Meat and offal. 194 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA11JA

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Surrendered

Authorised in:
Slovenia

Package description:
(ID3): 1 unspecified outer container with 12 Bottle (Glass) with 100 millilitre(s) (1200 millilitre(s))
(ID2): 1 unspecified outer container with 6 Bottle (Glass) with 100 millilitre(s) (600 millilitre(s))
(ID1): 1 unspecified outer container with 1 Bottle (Glass) with 100 millilitre(s) (100 millilitre(s))

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Bela-Pharm GmbH & Co. KG

Marketing authorisation date:
29/04/2019

Manufacturing sites for batch release:
Bela-Pharm GmbH & Co. KG

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
MR/V/0668/001

Date of authorisation status change:
27/09/2023

Reference member state:
Germany

Procedure number:
DE/V/0313/001

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents